07-023090.03 BASE AG

BADI 95,06,09

Act-E, 12(L3B) E(6-A2F, 7-A2E) L(3-D1D3)

*#P7:73.2.4 95.06.09 95DE-1020660 (95.12.17), CCTD 493 (4), Co9K (9.2), 10.3(,

Polymerisable chiral cpds., used a dopants in liq. crystal compsns. contg. condensable reactive gps., e.g. isocyanate, attached to a multivalent chiral gp., e.g. diranhydro-sorbityl, via spacer and mesogenic gps. (Ger)

C97-007478 R/BE CH DE FR 5B IT LINE MEYER F. SIEMENSMEYER K. ETZBACH K. Addnl. Data

SCHUHMACHER P 96 06 03 96EP-108872

Polymensable entral cpds of formula (I) are new

Z. (Y - A - Y - M - Y - N) in which A=a spacer, M=mesogenic gp: Y-Y=direct bond, O, S, COO, OCO, OCOO, CONR or NECO, <math>R=H or 1-4C all M=nvalent chiral gp: n = 2-n, at least one Z is a residue with a NCO, NCS, CNO, thirrane, azaridane, COOH, OH or ammo gp., and the otheristissate H or unreactive go-

USE

In electro-optical liq. crystal (LC) displays, as chiral dopants for

nematic or cholestone Lel compans, and as entral doponts for the production of coloured, reflecting, enolesteric LC (avers (claimed)

ADVANTAGE Provides claral depant, with a high twisting power capable of stable incorporation in cholesteric phases over a wide range of concuswiraout diffusing or erestallising out.

PREFERRED COMPOUNDS

-(T-Y4),-T-(la), in which T = n = 2, M = a gp. of formula divalent iso- or netero- cycloaliphatic gp- or iso- or netero-aromatic gp. Y' = , bridging gp as for Y', or -CH₂O₋, -OCH₂-, -CH=N- or -N=CH-, r = 0-3, pret, 0 or 1. T and Y can be the same or different when r = more than 0 or more than 1 - X = one of the following gps.as shown,

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in which L = 1-4C alkyl or alkovy, halogen, -COOR, -OCOR, -CONFIR or "NHCOR". At least one of the gps. Z-Y' = "CNO or "NCO (i.e. with Y' = direct bond). Pref., gps. Z, Y1, Y2, Y3, A, M, R and L are each identical.

PREPARATION

Cpds. (I) are prepri by known methods (eg. as described in DE-A 3917196) from chiral starting materials of formula X(OH)n, which are generally commercially available cpds. Gps. Z, A, M and X are pref. coupled by condensation reactions with the formation of bridging gps. (Y), e.g. by reacting a mesogenic carboxylate with a chiral OH cpd. to form an ester, or two OH cpds, to form an ether, etc.

A mixt of 4.5 g 4-isocyanatobenzoyl chloride, 3.5 g 2,5-bistrimethylsilyloxy-1,4,3,6-dianhydrosorbitol, 20 mg 4dimethylaminopyridine and 10 ml 1,2-dichlorobenzene was heated, at 165°C for 11 hrs. with removal of Me₁SiCl by distn. at 500 mbar. The mixt, was then worked up by cooling, filtration of the pptd, solid and recrystallisation, to give 2.5-bis-(4'-iso-cyanatobenzoyl)-1,4,3,6dianaydrosorbitol (1) in 83% yield. (13pp1712DwgNo.0/0)

SR EP399279 EP409066 EP564932 EP630892 US4631328 WO9516007

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